

Orbit & Sinus: What are you looking for?



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Objectives

- Recognize imaging findings in orbits & paranasal sinuses that will change patient management.
- Be able to develop “checklist” for imaging findings within orbits & paranasal sinuses that decreases likelihood of overlooking pertinent associated findings.

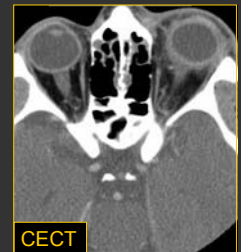
Orbits & Sinuses

- Infection
- Trauma
- Neoplasm



Orbit: Infectious

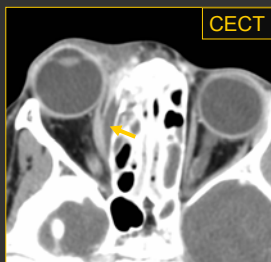
- **Pre - or post-septal**
- Most often secondary to underlying paranasal sinusitis
 - Maxillary & ethmoid most common
- Other etiologies:
 - Trauma
 - Bacteremia
 - Skin infections
 - Dental infections



Panophthalmitis

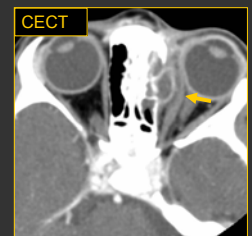
Orbit: Infectious

- Subperiosteal abscess
 - Spread from **sinus to orbit**
 - Transmission via valveless venous plexus
 - **Direct extension:**
 - Lamina papyracea dehiscence
 - Visual disturbance: 15-30%
 - Optic neuritis
 - Ischemia:
 - ↑ intraorbital pressure
 - Retinal ischemia: Central artery occlusion or thrombophlebitis



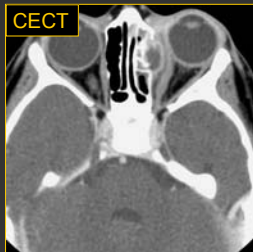
Subperiosteal Abscess: Orbit

- **CT:**
 - Medial orbital wall with adjacent sinusitis
 - Lentiform, rim enhancing
 - Medial rectus displacement
 - Lamina papyracea dehiscence
- **MR**
 - Post contrast: Rim enhancement; intra- & periorbital enhancement
 - **Fat suppression** optimal
- **Requires immediate attention!**
 - May result in blindness



Orbital Infection: Checklist

- ✓ Abscess?
- ✓ Evidence of cavernous sinus thrombosis?
 - Rare
- ✓ Evaluate brain – evidence of:
 - Meningitis? – Lumbar puncture
 - Subdural empyema?
 - Abscess?
 - Cerebritis?



Preseptal Cellulitis

- Anterior to orbital septum
- May be difficult to distinguish clinically from subperiosteal abscess
- Orbital involvement suspected:
 - Proptosis
 - Extraocular motility defects
 - Decreased vision

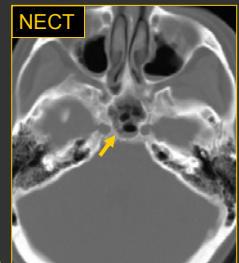


Preseptal Cellulitis

- In pediatric patients & in skilled hands → US useful for rapid evaluation of preseptal vs. postseptal involvement
- US limited in ability to assess:
 - Intracranial extension
 - Orbital apex
 - Paranasal sinuses

Sinus Infectious

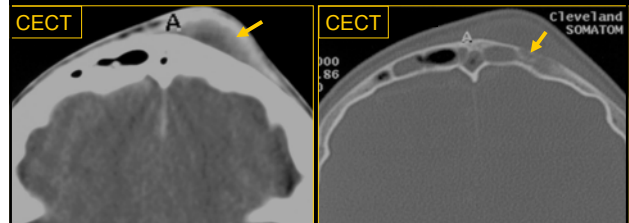
- Acute sinonasal inflammatory disease
 - Typically not imaged → Clinical diagnosis
 - Resolves with conservative measures
 - Imaging Features
 - Air-fluid level
 - Bubbly secretions
- Complicated sinonasal inflammatory disease → Image
 - Orbital &/or CNS complications



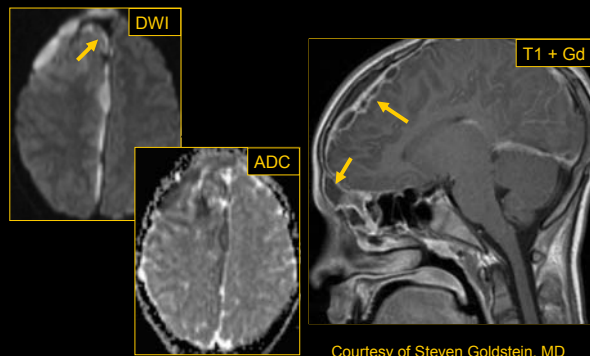
Acute sinusitis: Complications

- Local extension
 - Orbital: Sub-periosteal abscess
 - Intra-cranial : Empyema, meningitis, cerebritis, abscess
 - Superficial: Osteomyelitis, subgaleal abscess
- Venous occlusion: Cavernous sinus

Pott's Puffy Tumor

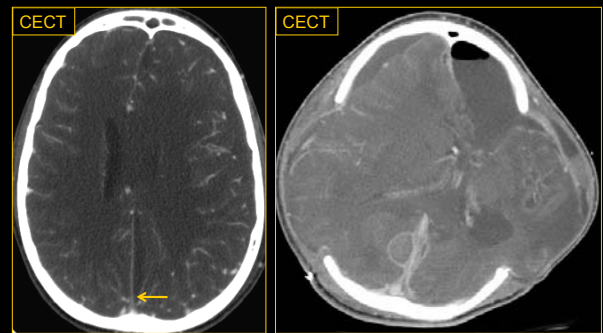


Subdural Empyema



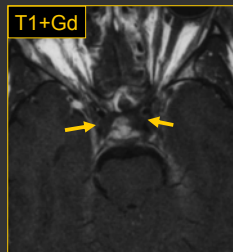
Courtesy of Steven Goldstein, MD

Subdural Empyema



Sinusitis: Checklist

- ✓ Evidence of local extension
 - ✓ Orbital
 - ✓ Intra-cranial
 - ✓ Superficial
- ✓ Evidence of venous occlusion



Fungal Sinusitis

- Subtypes:
 - Acute invasive
 - Chronic invasive
 - Chronic granulomatous invasive
 - Allergic fungal
 - Fungal mycetoma ("fungus ball")

Invasive
Non-invasive

Invasive = Presence of fungal hyphae within mucosa, submucosa, bone, or blood vessels of paranasal sinuses

Fungal Sinusitis: Acute Invasive

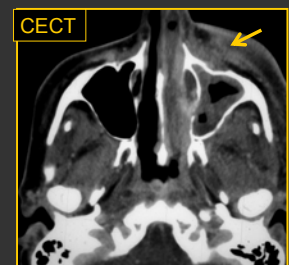
- Rapidly progressive
 - Mortality: 50-80%
- Diabetic or other immunocompromised
- Maxillary & ethmoid most common
- Adjacent bone erosion & soft tissue infiltration



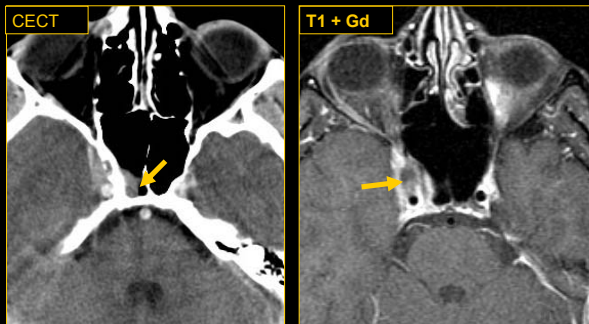
Mucor

Fungal Sinusitis: Acute Invasive

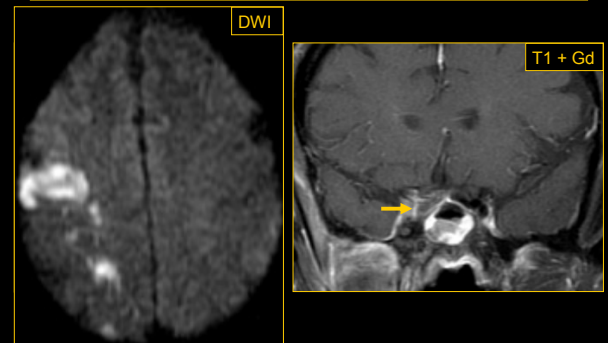
- CT/MR
 - Obliteration of periantral fat: **May be subtle!**
 - Late:
 - Intracranial/orbital spread
 - Leptomeningeal enhancement **may be subtle** in early stage
 - Bone destruction



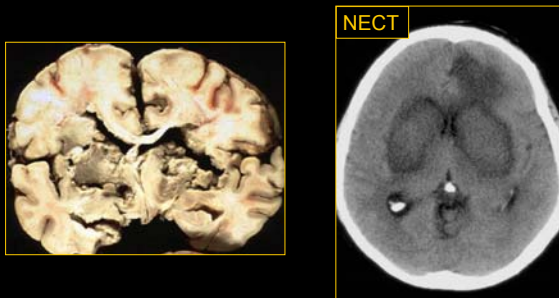
Beware Isolated Sphenoid Sinus Disease in Immunocompromised!



Mucormycosis

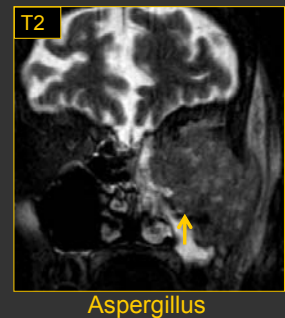


Mucor



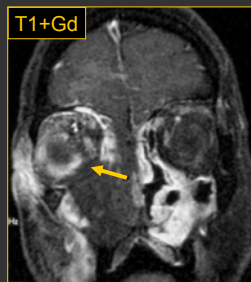
Invasive Fungal Sinusitis

- Mycetoma low T2
 - ? paramagnetic materials produced by fungi
 - ? semisolid, cheesy nature of mycetoma



Invasive Fungal Sinusitis Checklist

- ✓ Evaluate periantral fat
- ✓ Evaluate orbit & intracranially for involvement
- ✓ Evaluate cavernous sinus & internal carotid



Trauma: Orbit

- Eye trauma: 3% of ER visits
- Concomitant injury to brain, spinal cord, or facial bones common
- Injuries:
 - Intraorbital foreign body
 - Optic nerve
 - Open globe
- CT imaging study of choice

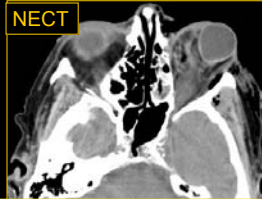


Globe Tenting

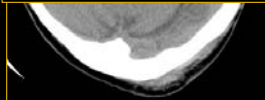
NECT



NECT



Emergent surgical decompression!!!

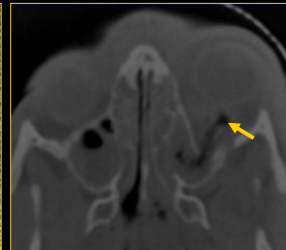
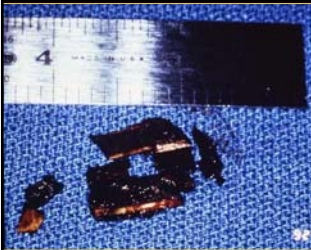


Orbital Foreign Bodies

- CT scan: Test of choice
- Most common cause: Hammering
- May be observed:
 - Smooth edges
 - Located in posterior orbit
- Removed:
 - Composed of **vegetable matter**
 - Iron containing can cause siderosis
 - Lead containing – may cause lead poisoning
 - Copper - sterile endophthalmitis
 - Easily accessible in anterior orbit

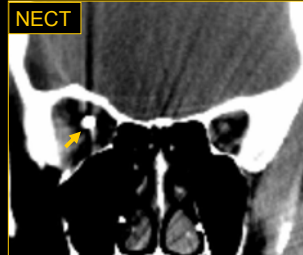


Foreign Body: Wood

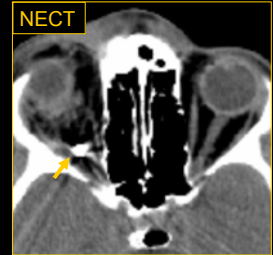


Orbital Foreign Body: "Double Perforation"

NECT



NECT



Trauma: Hemorrhagic choroidal detachment



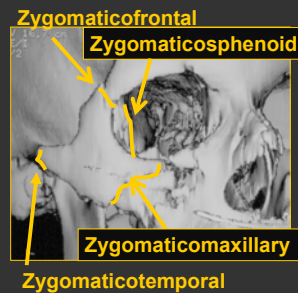
Orbital Trauma Checklist

- ✓ Presence of foreign body
- ✓ Hemorrhage
- ✓ Status of optic nerve
 - ✓ Orbital apex
- ✓ Intracranial injury



Sinus & Orbit: Zygomaticomaxillary Complex (ZMC) fracture

- Disjunction of zygoma from adjacent osseous connections
- “Tripod” abandoned
 - Quadripod



ZMC Fracture Complications/Checklist

- ✓ Infraorbital foramen – paresthesias
- ✓ Zygomatic arch impale coronoid process → trismus
- ✓ Fractured **uncinate** – posttraumatic sinus disease
- ✓ **Lateral rectus** impaled by lateral wall of orbit
- ✓ Ruptured Globe



Neoplastic

- Orbit
 - **Retinoblastoma**
 - **Melanoma**
 - Lymphoma
 - Metastasis
 - Rhabdomyosarcoma
- Sinus
 - **SCC**
 - **Adenocarcinoma**
 - **Esthesioneuroblastoma**
 - **Inverting papilloma**
 - **SNUC**
 - Lymphoma
 - Melanoma



Melanoma

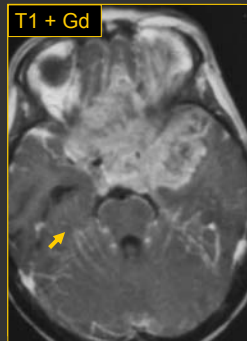
Retinoblastoma

- Primary retinal malignant neoplasm
- 90-95% before age 5
 - **Most common** intraocular tumor of childhood
 - Hereditary form earlier
- Location:
 - **Unilateral**: 70-75%; 30% multifocal
 - **Bilateral**: 25-30%
 - **Trilateral** (midline neuroblastic tumor + bilateral ocular): Rare
 - **Quadrilateral** (suprasellar + bilateral ocular + pineal): Rare



Retinoblastoma

- Ophthalmoscopic diagnosis primarily
 - Small gray-white intraretinal lesions, calcification, seeding
 - Ultrasonography: 80% accurate
- Staging:
 - Stage 1: Confined to globe
 - Stage 2: Extraocular extension to orbit or optic nerve
 - Stage 3: Extra-orbital extension
- 92% 5-year survival for intra-ocular lesions

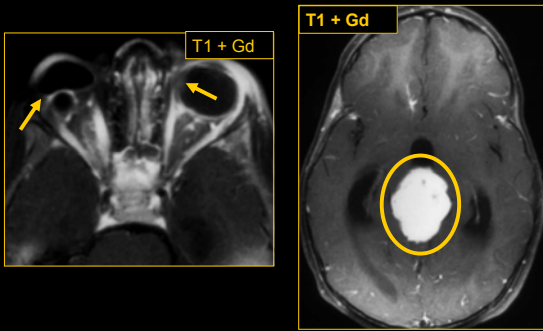


Retinoblastoma: Imaging

- CT: Calcified intraocular mass **90-95%**
 - Utilize thin section (1.5mm)
- MRI
 - ↑T1, ↓T2
 - Optic nerve & transscleral extension?
 - Anterior segment involvement?

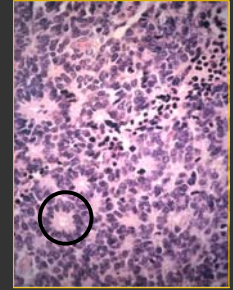


Trilateral Retinoblastoma



Retinoblastoma: Pathology

- **Neuroectodermal** origin: primitive embryonal retinal cells (retinoblasts)
- **Rosettes:** Flexner-Wintersteiner
- Highly malignant: Necrosis, mitotic figures
- Calcification

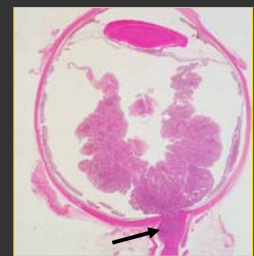
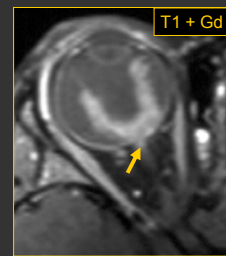


Retinoblastoma

- **90% cure rate** for non-invasive
- Biopsy carries risk of seeding – radiologic diagnosis critical
- Regular **screening** for children with family history
- Surveillance **through age 7** years assess for development of metachronous disease



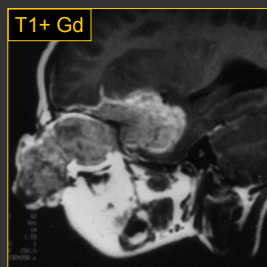
Retinoblastoma



- Optic nerve/intraorbital extension: 10-15%
– Poor prognostic factor

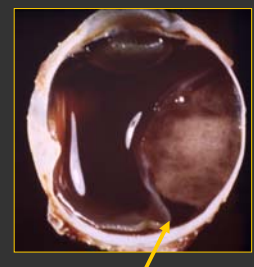
Retinoblastoma: Checklist

- ✓ Intracranial **trilateral** or **quadrilateral** disease?
- ✓ Involvement of anterior segment?
- ✓ Optic nerve involvement or transscleral extension?



Melanoma

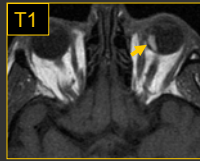
- Melanoma: Most common primary intraocular tumor in adults
- Arises from melanocytes within the choroid
- Whites (15:1)
– Incidence increases with age
- Metastasize to liver & lung



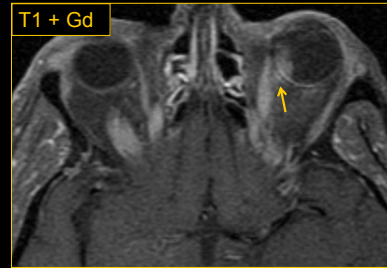
Retinal detachment

Melanoma: Imaging

- CT:
 - High density
 - Enhance
- MRI:
 - T1 hyperintense
 - T2 hypointense
 - **Amelanotic**: T1 hypointense, T2 hyperintense
 - Lesions elevated > 3 mm usually seen on MR
 - < 3mm better evaluated by US



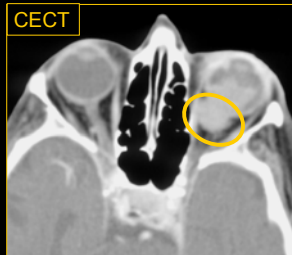
Extraocular Invasion



Poorer prognosis & different therapeutic implications

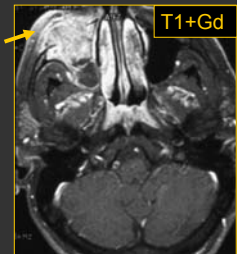
Melanoma Checklist

✓ Extraocular invasion



Sinonasal Neoplasms

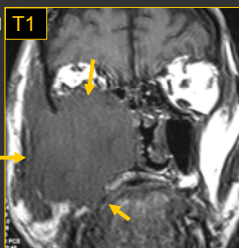
- SCCa
- Esthesioneuroblastoma
- Adenocarcinoma
- Sinonasal undifferentiated carcinoma (SNUC)
- Lymphoma
- Melanoma



SCCa

Squamous Cell Carcinoma

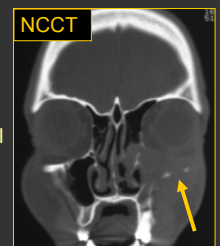
- Malignant epithelial tumor
 - Most common malignancy of sinonasal area
- Maxillary **antrum** most common (85%)
- Pre-surgical evaluation of extent:
 - Anterior: SQ tissue of cheek
 - Superior: Orbit
 - Inferior: Hard palate, maxillary alveolar ridge
 - Posterior: Retroantral fat & PPF
 - Perineural spread



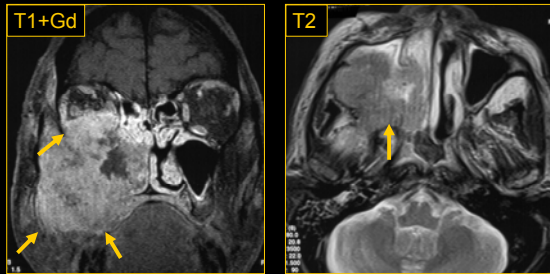
Lymph node: 15% at presentation – retropharyngeal & jugulodigastric

SCC: Imaging

- CT:
 - Bone destruction
 - Heterogeneous enhancement
- MRI
 - T1: Intermediate signal
 - T2: Lower signal than most sinonasal malignancies
 - Post contrast: Heterogeneously enhances
 - Enhancement < adenocarcinoma, esthesioneuroblastoma, melanoma
 - Fat sat: **Perineural spread**

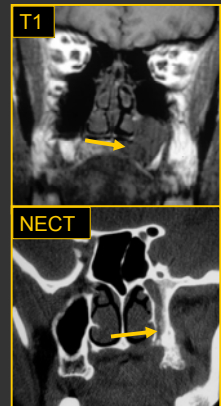


SCC



Perineural Spread

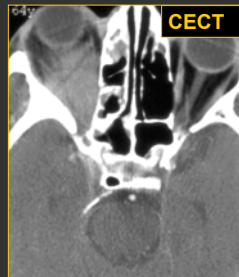
- Perineural spread
 - Widened foramen or canal
 - Enlarged enhancing nerve
 - Obliteration of fat at skull base foramen



Adenocarcinoma

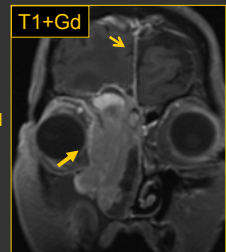
Neoplasm: Checklist

- ✓ Evaluate for extension: Orbit, palate
- ✓ Evaluate pterygopalatine fossa and for evidence of perineural spread
- ✓ Malignant adenopathy



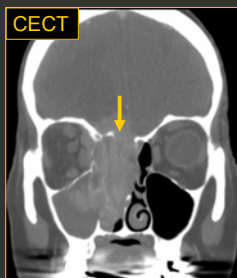
Esthesioneuroblastoma

- Neuroendocrine malignancy of neural crest origin
- Arises from olfactory epithelium
- Bimodal age distribution: 2nd & 6th decades
- Malignant cervical lymph nodes at presentation: 20%
- Long term follow up: Tend to recur late

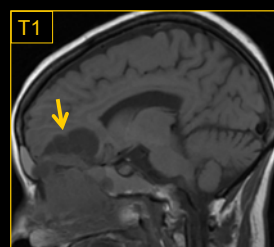


Esthesioneuroblastoma

- CT
 - Bone remodeling mixed with destruction
 - Homogeneously enhances – may have areas of necrosis
- MRI
 - T1: Hypo- to isointense
 - T2: Iso- to hyperintense



Esthesioneuroblastoma



Intracranial peripheral tumor cysts suggestive of diagnosis

Esthesioneuroblastoma: Checklist

- ✓ Degree of spread
- ✓ Lymph node or distant metastasis



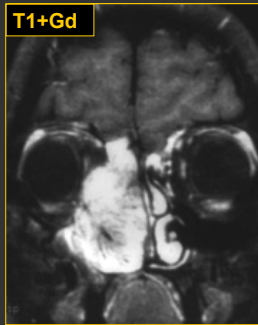
Inverted Papilloma

- Epithelial tumor of nasal mucosa
- Most commonly originates in lateral wall of nose
- Spread into adjacent sinuses & possibly orbit & CNS
- Morphology: “Cerebriform”
- 5-15% Associated with SCCa



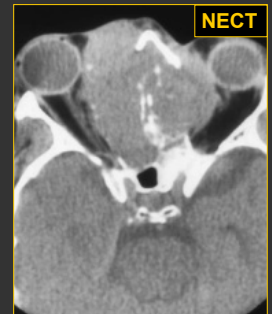
Inverted Papilloma

- High recurrence rate & associated SCCa → Imaging follow up
- Identifying adjacent areas of invasion may alter surgical approach
- Locally invasive disease difficult to evaluate on nasal endoscopy

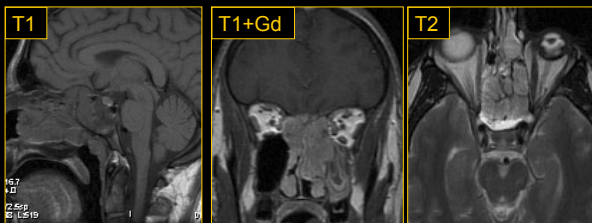


Inverted Papilloma

- CT:
 - Large remodels nasal cavity & invades/obstructs adjacent sinuses
 - Osseous destruction – Consider associated SCCa



Inverted Papilloma



- MR:
 - T2: Curvilinear striations → cerebriform
 - Enhance: May have convoluted appearance
 - If appears invasive consider SCCa

Inverted Papilloma: Checklist

- ✓ Adjacent areas of invasion on imaging
- ✓ Association with SCCa

Conclusion

- Developing a “checklist” for imaging findings within orbits & paranasal sinuses decreases likelihood of overlooking pertinent associated findings.



Courtesy Steven Goldstein,
MD